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(54) Title: CONTRAST MEDIA FOR ULTRASONIC IMAGING

(57) Abstract

Novel contrast media for use in ultrasonic imaging are described. Such contrast media may be comprised of an aqueous solution of one or more biocompatible polymers, wherein said biocompatible polymers are coated with and/or in admixture with at least one silicon containing compound. Alternatively, the contrast media may be comprised of an aqueous solution of one or more biocompatible synthetic polymers, or an aqueous solution of cellulose. The contrast media may be employed, if desired, with anti-gas agents and/or suspending agents.

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TITLECONTRAST MEDIA FOR ULTRASONIC IMAGINGBACKGROUND OF THE INVENTIONField of the Invention

5 This invention relates to the field of ultrasonic imaging, and more specifically to the use of polymers as contrast agents for ultrasonic imaging.

Description of the Prior Art

10 There are a variety of imaging techniques that have been used to diagnose disease in humans. One of the first imaging techniques employed was X-rays. In X-rays, the images produced of the patient's body reflect the different densities of body structures. To improve the diagnostic utility of this imaging technique, contrast
15 agents are employed in an attempt to increase the differences in density between various structures, such as between the gastrointestinal tract and its surrounding tissues. Barium and iodinated contrast material, for example, are used extensively for X-ray gastrointestinal studies to visualize the esophagus, stomach, intestines and rectum. Likewise, these contrast agents are used for X-ray computed tomographic studies to improve visualization
20 of the gastrointestinal tract and to provide, for example, contrast between the tract and the structures adjacent to it, such as the vessels or the lymph nodes. Such gastrointestinal contrast agents permit one to increase the density inside the esophagus, stomach, intestines and

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rectum, and allow differentiation of the gastrointestinal system from surrounding structures.

- Ultrasound is a more recent imaging technique which, unlike X-rays, does not utilize ionizing radiation.
- 5 Instead, in ultrasound, sound waves are transmitted into a patient. These sound waves are then reflected from tissue in the patient and are received and processed to form an image. Since ultrasound does not employ ionizing radiation to produce these images, ultrasound is less invasive and
- 10 safer to the patient than X-ray imaging techniques. Ultrasound, however, suffers at times in imaging clarity in comparison to X-rays, particularly where imaging of the gastrointestinal tract is involved. In ultrasound, one major problem is the presence of air/fluid interfaces,
- 15 which results in shadowing of the ultrasound beam. Shadowing, in turn, prevents the ultrasound beam from penetrating beyond the air/fluid interface, and thus prevents visualization of structures near any air pockets. Another problem with ultrasound is the difficulty in
- 20 imaging adjacent hypoechoic structures, that is, structures that are only minimally reflective of the ultrasound beam and, therefore, result in a low ultrasound signal. These problems are particularly evident in the gastrointestinal region, with its many air/liquid interfaces and its
- 25 adjacent fluid-filled mucosal lining and often fluid-containing lumen. If better contrast agents were available for ultrasound, the diagnostic accuracy and the overall usefulness of ultrasound as an imaging modality, particularly in the gastrointestinal region, would be
- 30 greatly enhanced.

In the past, investigators have attempted to solve the problems associated with gastrointestinal ultrasonic imaging by using water to fill the gastrointestinal tract. Water, however, was found to simply mix with the gas, and thus much of the shadowing resulting from the presence of air/fluid interfaces remained. In addition, the fact that water is absorbed by

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the bowel decreased its ability to serve any meaningful contrast enhancement function distally within the tract. Furthermore, the water is hypoechoic, and its presence adjacent to the fluid-filled hypoechoic mucosal lining of 5 this region resulted in little differentiation of the tract lumen and from its lining. Intravenously administered glucagon has been employed in connection with such gastrointestinal imaging, since glucagon administered in this fashion relaxes the bowel by decreasing peristalsis. 10 Although a helpful ultrasound adjunct, this, however, does not address such problems as shadowing caused by air/fluid interfaces and low differentiation caused by the presence of adjacent hypoechoic structures.

The need is great for contrast agents useful in 15 ultrasonic imaging of various regions of the body, particularly those useful in imaging the gastrointestinal tract. The present invention is directed to these important ends.

SUMMARY OF THE INVENTION

20 The present invention is directed to a contrast medium useful for ultrasonic imaging.

Specifically, the invention pertains to a contrast medium comprising an aqueous solution of at least one biocompatible polymer, wherein said biocompatible 25 polymer is coated with and/or in admixture with at least one silicon containing compound.

The invention also pertains to a contrast medium comprising an aqueous solution of at least one biocompatible synthetic polymer.

30 The invention further pertains to a contrast medium comprising an aqueous solution of cellulose.

Further, the subject invention encompasses a method of providing an image of an internal region of a patient, especially an image of the gastrointestinal region 35 of the patient, said method comprising (i) administering to the patient one or more of the aforementioned contrast

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media, and (ii) scanning the patient using ultrasonic imaging to obtain visible images of the region.

Still further, the present invention comprises a method for diagnosing the presence of diseased tissue in a patient, especially in the gastrointestinal region of the patient, said method comprising (i) administering to the patient one or more of the foregoing contrast media, and (ii) scanning the patient using ultrasonic imaging.

Finally, the present invention contemplates kits including the foregoing contrast media.

The present ultrasound contrast agents are particularly useful when employed in the gastrointestinal region, serving to improve visualization by displacing gas within the tract and providing contrast to the bowel lumen by filling it with material which has an echogenicity different from the adjacent mucosa.

These and other aspects of the invention will become more apparent from the following detailed description when taken in conjunction with the following drawings.

BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 is a diagrammatic view of a portion of the gastrointestinal tract of a patient showing representative air and fluid levels and air bubbles naturally present in the stomach and small intestines.

Fig. 2A is a diagrammatic view of a typical ultrasound image of the gastrointestinal tract of Fig. 1, showing the shadowing that results from air bubbles and air/fluid levels.

Fig. 2B is a typical photograph of an ultrasound image of the gastrointestinal tract of Fig. 1, showing the shadowing that results in the stomach from air bubbles and air/fluid levels.

Fig. 2C is a typical photograph of an ultrasound image of the gastrointestinal tract of Fig. 1, showing the shadowing that results in a portion of the small intestine from air bubbles and air/fluid levels.

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Fig. 3A is a diagrammatic view of an ultrasound image of the gastrointestinal tract of Fig. 1 after consumption of 750 ml of a 5% cellulose and 0.25% xanthan gum solution of the invention. Significant air/fluid levels and air bubbles are no longer present. Improved visualization of the gastrointestinal tract in general and improved visualization of the mucosa as distinguished from the tract lumen is observed.

Fig. 3B is a photograph of an ultrasound image of the gastrointestinal tract of Fig. 1 after consumption of 750 ml of a 5% cellulose and 0.25% xanthan gum solution of the invention. Improved visualization of the gastrointestinal tract in general and improved visualization of the mucosa as distinguished from the tract lumen is observed. In Fig. 3B, S denotes stomach, D denotes duodenum, and P denotes pancreas.

Fig. 4 is a graph showing the dB reflectivity of cellulose coated with silicone and uncoated cellulose in vitro. The coated cellulose exhibits a significantly better dB reflectivity.

Fig. 5 is a graph showing the dB reflectivity of cellulose fibers 18 μ in length coated with varying amounts of silicone and simethicone in vitro.

Fig. 6 is a graph showing the dB reflectivity of uncoated cellulose fibers of varying length at different concentrations of cellulose and different transducer frequencies in vitro.

Fig. 7 is a graph showing the dB reflectivity of uncoated cellulose fibers of varying length at different concentrations of cellulose and different transducer frequencies in vitro.

Fig. 8 is graph showing the dB reflectivity of cellulose fibers of varying length coated with 1% of differnt silicon compounds in vitro.

DETAILED DESCRIPTION OF THE INVENTION

Any of the wide variety of biocompatible polymers known in the art may be employed in the medium and methods

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of the subject invention. The term biocompatible, used herein in conjunction with the term polymers, is employed in its conventional sense, that is, to denote polymers that do not substantially interact with the tissues, fluids and 5 other components of the body in an adverse fashion in the particular application of interest. As will be readily apparent to those skilled in the art, there are numerous types of such polymers available.

The polymers useful in the present invention can 10 be of either natural, semisynthetic or synthetic origin. As used herein, the term polymer denotes a compound comprised of two or more repeating monomeric units, and preferably 10 or more repeating monomeric units. The term semisynthetic polymer, as employed herein, denotes a natural 15 polymer that has been chemically modified in some fashion. Exemplary natural polymers suitable for use in the present invention include naturally occurring polysaccharides. Such polysaccharides include, for example, arabinans, fructans, fucans, galactans, galacturonans, glucans, 20 mannans, xylans (such as, for example, inulin), levan, fucoidan, carragheenan, galactocarolose, pectic acid, amylose, pullulan, glycogen, amylopectin, cellulose, dextran, pustulan, chitin, agarose, keratan, chondroitan, 25 dermatan, hyaluronic acid, alginic acid, xanthan gum, starch and various other natural homopolymer or heteropolymers such as those containing one or more of the following aldoses, ketoses, acids or amines: erythrose, threose, ribose, arabinose, xylose, lyxose, allose, altrose, glucose, mannose, gulose, idose, galactose, 30 talose, erythrulose, ribulose, xylulose, psicose, fructose, sorbose, tagatose, glucuronic acid, gluconic acid, glucaric acid, galacturonic acid, mannuronic acid, glucosamine, galactosamine, and neuraminic acid, and naturally occurring derivatives thereof. Exemplary semisynthetic polymers 35 include carboxymethylcellulose, hydroxymethylcellulose, hydroxypropylmethylcellulose, methylcellulose, and methoxycellulose. Exemplary synthetic polymers suitable

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for use in the present invention include polyethylenes (such as, for example, polyethylene glycol, polyoxyethylene, and polyethylene terephthalate), polypropylenes (such as, for example, polypropylene glycol), polyurethanes (such as, for example, polyether polyurethane ureas), pluronic acids and alcohols, polyvinyls (such as, for example, polyvinyl alcohol, polyvinylchloride and polyvinylpyrrolidone), nylon, polystyrene, polylactic acids, fluorinated hydrocarbons, fluorinated carbons (such as, for example, polytetrafluoroethylene), and polymethylmethacrylate, and derivatives thereof.

Preferably, the polymer employed is one which has a relatively high water binding capacity. When used, for example, in the gastrointestinal region, a polymer with a high water binding capacity binds a large amount of free water, enabling the polymer to carry a large volume of liquid through the gastrointestinal tract, thereby filling and distending the tract. The filled and distended gastrointestinal tract permits a clearer picture of the region. In addition, where imaging of the gastrointestinal region is desired, preferably the polymer employed is also one which is not substantially degraded within and absorbed from the gastrointestinal region. Minimization of metabolism and absorption within the gastrointestinal tract is preferable, so as to avoid the removal of the contrast agent from the tract as well as avoid the formation of gas within the tract as a result of this degradation. Moreover, particularly where gastrointestinal usage is contemplated, the polymers are preferably such that they are capable of displacing air and minimizing the formation of large air bubbles within the polymer composition.

In accordance with the invention, the polymers may be coated with and/or be in admixture with a silicon containing compound. As used herein, the phrase silicon containing compound denotes both organic and inorganic compounds containing the element silicon (Si). By

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- adixture, it is meant that the silicon containing compound is simply added to the polymer containing medium, and is not chemically bound to the polymer. By the term coated, it is meant that the silicon containing compound is
- 5 chemically bound to the polymer. Suitable silicon containing compounds include silicone, simethicones (such as simethicone and protected simethicone, the latter compound being polydimethylsiloxane with 4% to 4.5% silicon dioxide), siloxanes (such as polysiloxane, polydimethyl-
10 siloxane, polymethylvinylsiloxane, polymethylphenol-siloxane, polydiphenylsiloxane, octamethylcyclotetrasiloxane and siloxane glycol polymers), silanes (such as gamma-aminopropyltriethoxysilane, diethylaminomethyltriethoxysilane), silicon dioxide, siloxyalkylene polymers,
15 linear or cyclic silazanes (such as hexadimethylsilazane, hexamethyldisilazane, hexaphenylcyclotrisilazane and octamethylcyclotetrasilazane), silyl compounds (such as N'N'-bis-(trimethylsilyl)acetamide and N-(dimethyl(gamma-cyanopropyl)-silyl)-N-methylacetamide), and siliceous earth. Other suitable silicon compounds will be readily apparent to those skilled in the art, and are described, for example, in Hertl et al., J. Phys. Chem., Vol. 75, No. 14, pp. 2182-2185 (1971). Such silicon compounds may be easily prepared using conventional chemical synthesis
20 methodology, such as is described in Chem. Abstracts, Vol. 69, No. 61770x (1968), Vol. 70, No. 1164505k (1969), Vol. 74, No. 59904u (1971), Vol. 75, No. 9271v (1971), and Vol. 77, No. 37150q (1972), or may be obtained from various commercial sources. The silicon containing compounds
25 surprisingly serve a number of important functions in the contrast medium, as both coating or admixed with the polymers, assisting in decreasing surface tension, minimizing foaming, increasing reflectivity, and/or lessening acoustic attenuation.
- 30 To prepare the biocompatible polymer coated with the silicon containing compound, a polymer reactive with the desired silicon containing compound is mixed therewith
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under conditions suitable for chemical bonding of the polymer and the silicon compound. Such reactive polymers and suitable conditions will be readily apparent to one skilled in the art, once in possession of the present disclosure. Suitable procedures include those described in Noll, Chemistry and Technology of Silicones, pp. 515-521 (Academic Press 1968), Hertl et al., *J. Phys. Chem.*, Vol. 75, No. 14, pp. 2182-2185 (1971), and Stark et al., *J. Phys. Chem.*, Vol. 72, No. 8, pp. 2750-2754 (1968), the disclosures of each of which are incorporated herein by reference in their entirety. Particularly reactive with many of the silicon containing compounds are polymers containing free hydroxyl groups, such as cellulose and polyethylene glycol.

Cellulose, a naturally occurring polymer that exhibits a high water binding capacity, when coated or admixed with a silicon containing compound, is particularly preferred for use in the subject invention, especially when gastrointestinal usage is desired, resulting in excellent echogenicity when ultrasound is applied. As a result of their high water binding capacity, the cellulosic compounds pass through the gastrointestinal tract in a liquid medium distending the tract and displacing gas within the tract. In addition, since cellulose is not degraded and absorbed within the tract, it does not result in the formation of any gases and is not removed from the tract along its route. Furthermore, cellulose is highly effective in displacing air and avoiding the formation of large air bubbles within the polymer composition.

Polyvinylpyrrolidone, polyethylene glycol, and other polyethylenes, when coated or admixed with a silicon containing compound, are also preferable polymers, particularly where gastrointestinal imaging is contemplated, functioning in a fashion similar to cellulose in binding water and distending the gastrointestinal tract, and resulting in highly improved visualization of the bowel and adjacent structures by ultrasound. Polyethylene glycol

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and other polyethylenes are also very effective in preventing the formation of and in dispersing large gas bubbles. The silicon containing compounds employed with the foregoing polymers serve to decrease surface tension (thus allowing gas in the solution to escape more easily), minimize foaming, increase reflectivity, and/or lessen acoustic attenuation in the resultant contrast media.

5 The polymers of the present invention may be employed as solids in various shapes and forms, such as, for example, fibers, beads, and the like. As those skilled in the art will recognize, the size of the individual polymer fibers, beads, etc., can also vary widely. Preferably, however, the length of any polymer fibers and diameter of any polymer beads is between about 0.1 and 10 about 200 microns, more preferably between about 5 and 15 about 100 microns, most preferably between about 10 and about 20 microns. Fibers are preferred, and in general, shorter fiber lengths have been found to possess better 20 acoustic properties and have better suspension uniformity than longer fiber lengths.

If desired, the polymers employed in the invention may be in liquid form, that is, liquid at physiological temperatures. The liquid polymer has its own inherent unique echogenicity which allows differentiation 25 of bodily structures. Polyethylene glycol (PEG) of low molecular weight functions well in this regard (the lower molecular weight polymers being liquid at physiological temperatures) as do fluorinated hydrocarbons. As one skilled in the art will recognize, there are many polymers 30 which are liquid at physiological temperatures and can be employed in this fashion.

The polymers of the invention may be employed in an aqueous solution as a contrast medium for ultrasound imaging.

35 If desired, in addition, the polymer solutions may be employed in conjunction with an additional anti-gas agent. As used herein the term anti-gas agent is a

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compound that serves to minimize or decrease gas formation, dispersion and/or adsorption. A number of such agents are available, including antacids, antiflatulents and antifoaming agents. Such antacids and antiflatulents 5 include, for example, activated charcoal, aluminum carbonate, aluminum hydroxide, aluminum phosphate, calcium carbonate, dihydroxyaluminum sodium carbonate, magaldrate magnesium oxide, magnesium trisilicate, sodium carbonate, loperamide hydrochloride, diphenoxylate, hydrochloride with 10 atropine sulfate, Kaopectate™ (kaolin) and bismuth salts. Suitable antifoaming agents useful as anti-gas agents include polyoxypropylene/polyoxyethylene copolymers, polyoxyalkylene amines and imines, branched polyamines, mixed oxyalkylated alcohols, sucroglycamides (celynols), 15 polyoxylalkylated natural oils, halogenated silicon-containing cyclic acetals, lauryl sulfates, 2-lactylic acid esters of unicarboxylic acids, triglyceride oils. Particles of polyvinyl chloride may also function as anti-foaming agents in the subject invention. Of course, as 20 those skilled in the art will recognize, any anti-gas agents employed must be suitable for use within the particular biological system of the patient in which it is to be used.

The polymer solutions may also, if desired, be 25 employed with a suspending or viscosity increasing agent, referred to herein collectively as a suspending agent. The phrase suspending agent, as used herein, denotes a compound that assists in providing a relatively uniform or homogeneous suspension of polymer through out the aqueous 30 solution. A number of such agents are available, including xanthum gum, acacia, agar, alginic acid, aluminum monostearate, unpurified bentonite, purified bentonite, bentonite magma, carbomer 934P, carboxymethylcellulose calcium, carboxymethylcellulose sodium, carboxymethyl- 35 cellulose sodium 12, carrageenan, cellulose (microcrystalline), carboxymethylcellulose sodium, dextrin, gelatin, guar gum, hydroxyethyl cellulose, hydroxypropyl

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cellulose, hydroxypropyl methylcellulose, magnesium aluminum silicate, methylcellulose, pectin, polyethylene oxide, polyvinyl alcohol, povidone, propylene glycol alginate, silicon dioxide, silicon dioxide colloidal, 5 sodium alginate, and tragacanth.

Wide variations in the amounts of the polymer, silicon containing compound, anti-gas agent, and suspending agent can be employed in the contrast medium of the invention. Preferably, however, the polymer, when employed 10 in solid form, is present in an aqueous solution in a concentration of at least about 0.2% by weight, more preferably at least about 0.5% by weight, even more preferably at least about 1% by weight. Of course, as those skilled in the art would recognize, within these 15 parameters, the optimum polymer concentration will be influenced by the molecular weight of the polymer, its water binding capacity, its particular echogenicity, as well as other characteristics of the particular polymer employed. In the case of cellulose, for example, the 20 polymer is most preferably present in a concentration of between about 1% and about 5% cellulose by weight. When a medium to high molecular weight polyethylene glycol or polyvinylpyrrolidone is used as the polymer, the concentration is most preferably between about 1% to about 25 15% by weight. When the polymers are in liquid form, generally the polymer is present in a somewhat higher concentration, such as preferably in a concentration of at least about 10% by weight, more preferably at least about 20% by weight, even more preferably at least about 30% by 30 weight. For example, when liquid polyethelene glycol is employed, the concentration is most preferably between about 10% and about 90% by weight. Similarly, when a liquid perfluorocarbon is used, the concentration is most preferably between about 1% and about 90% by weight. With 35 respect to the silicon containing compound, preferably the concentration is between about 0.1% by weight and about 20% by weight, more preferably between about 0.5% by weight and

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about 10% by weight, most preferably between about 1% by weight and about 5% by weight, whether present as a coating (that is, chemically bound to the polymer) or as an admixture (that is, added to the polymer medium but not chemically bound to the polymer). Generally, because of an increased ability to remain in a homogeneous suspension, the lower amounts of silicon containing compound are preferred.

In a preferably embodiment, the contrast medium of the invention is degassed, most preferably by either autoclaving the contrast medium, or by sonicating the contrast medium under vacuum (processes which forces gas out of solution as well as sterilizes the product), and then bottling the contrast medium under vacuum. The vacuum bottling is used so as to keep any new gas bubbles from forming in the product. The degassing and vacuum bottling serves to enhance the ultimate ecogenicity and minimize shadowing when the contrast medium is employed *in vivo*.

If desired, an antimicrobial agent may be included in the contrast medium to prevent bacterial overgrowth in the medium. Antimicrobial agents which may be employed include benzalkonium chloride, benzethonium chloride, benzoic acid, benzyl alcohol, butylparaben, cetylpyridinium chloride, chlorobutanol, chlorocresol, cresol, dehydroacetic acid, ethylparaben, methylparaben, methylparaben sodium, phenol, phenylethyl alcohol, phenylmercuric acetate, phenylmercuric nitrate, potassium benzoate, potassium sorbate, propylparaben, propylparaben sodium, sodium benzoate, sodium dehydroacetate, sodium propionate, sorbic acid, thimerosal, and thymol.

The present invention is useful in imaging a patient generally, and/or in specifically diagnosing the presence of diseased tissue in a patient. The imaging process of the present invention may be carried out by administering a contrast medium of the invention to a patient, and then scanning the patient using ultrasound imaging to obtain visible images of an internal region of a

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patient and/or of any diseased tissue in that region. By region of a patient, it is meant the whole patient, or a particular area or portion of the patient. The contrast medium is particularly useful in providing images of the 5 gastrointestinal region, but can also be employed more broadly such as in imaging any body cavities, or in other ways as will be readily apparent to those skilled in the art, such as in imaging the vasculature, liver, and spleen, and for use in tissue characterization. The phrase 10 gastrointestinal region or gastrointestinal tract, as used herein, includes the region of a patient defined by the esophagus, stomach, small and large intestines and rectum. By body cavities it is meant any region of the body having an open passage, either directly or indirectly, to the 15 external environment, such regions including the gastrointestinal region, the sinus tract, the fallopian tubes, etc. The patient can be any type of mammal, but most preferably is a human. Any of the various types of ultrasound imaging devices can be employed in the practice 20 of the invention, the particular type or model of the device not being critical to the method of the invention.

As one skilled in the art would recognize, administration of the contrast medium to the patient may be carried out in various fashions, such as orally, rectally 25 or by injection. When the region to be scanned is the gastrointestinal region, administration of the contrast medium of the invention is preferably carried out orally or rectally. When other body cavities such as the fallopian tubes or sinus tracts are to be scanned, administration is 30 preferably by injection. The useful dosage to be administered and the particular mode of administration will vary depending upon the age, weight and the particular mammal and region thereof to be scanned, and the particular contrast medium of the invention to be employed.

35 Typically, dosage is initiated at lower levels and increased until the desired contrast enhancement is achieved. By way of guidance, for gastrointestinal usage,

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about 10 ml of contrast medium per kg weight of the patient is administered (that is, a 70 kg patient would be administered about 700 ml of contrast medium). Various combinations of biocompatible polymers, silicon containing compounds, anti-gas agents, suspending agents, and other agents may be used to modify the echogenicity or ultrasonic reflectance of the gastrointestinal contrast agent, as well as to achieve the desired viscosity, gastric transit time, osmolality and palatability.

In carrying out the method of the present invention, the contrast medium can be used alone, or in combination with other diagnostic, therapeutic or other agents. Such other agents include flavoring or coloring materials, antioxidants, anticaking agents (to prevent caking on settling), tonicity agents (to optionally adjust osmolality to be iso-osmotic), and wetting agents (to facilitate mixing of the components in the contrast medium). Flavoring materials include, for example, aspartame, dextrates, dextrose, dextrose excipient, fructose, mannitol, saccharin, saccharin calcium, saccharin sodium, sorbitol, sucrose, sugar (compressible), and sugar (confectioner's syrup). Coloring materials include, for example, caramel and ferric oxide (e.g., red, yellow, black, or blends thereof). Antioxidants include, for example, ascorbic acid, ascorbyl palmitate, butylated hydroxyanisole, butylated hydroxytoluene, hypophosphorous acid, monothioglycerol, propyl gallate, sodium formaldehyde sulfoxylate, sodium metabisulfite, sodium thiosulfate, sulfur dioxide tocopherol, and tocopherols excipient. Anticaking agents include, for example, calcium silicate, magnesium silicate, and silicon dioxide (either colloidal or talc). Tonicity agents include, for example, dextrose, glycerin, mannitol, potassium chloride, sodium chloride, and propylene glycol. Wetting agents include, for example, benzalkonium chloride, benzethonium chloride, cetylpyridinium chloride, docusate sodium, nonoxynol 9, nonoxynol 10, octoxynol 9, polyoxamer, poloxyl 35 castor

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oil, polyoxyl 40 hydrogenated castor oil, polyoxyl 50 stearate, polyoxyl 10 oleyl ether, polyoxyl 20 cetostearyl ether, polyoxyl 40 stearate, polysorbate 20, polysorbate 40, polysorbate 60, polysorbate 80, sodium lauryl sulfate, 5 sorbitan monolaurate, sorbitan monooleate, sorbitan monopalmitate, sorbitan monostearate, and tyloxapol. The foregoing additional agents are conventional, and are described, for example, in the U.S. Pharmacopeia National Formulary, 22nd Revision, January 1, 1990, Mack Printing Company, Easton, PA.

The media of the present invention have been shown to be extremely useful as contrast enhancement agents in ultrasonic imaging, particularly in imaging of the gastrointestinal region.

15 Kits useful for ultrasonic imaging in accordance with the present invention comprise a contrast medium of the present invention, that is, an aqueous solution of at least one biocompatible polymer, wherein said biocompatible polymer is coated with and/or in admixture with at least 20 one silicon containing compound, in addition to conventional ultrasonic imaging kit components. Such conventional ultrasonic imaging kit components are well known, and include, for example, anti-gas agents, suspending agents, flavoring materials, coloring agents and 25 antioxidants, as described above.

A particularly preferred formulation of the contrast medium of the present invention useful in ultrasonic imaging, particularly of the gastrointestinal region, is a degassed solution comprised of the following 30 components:

- (i) 0.5% by weight of an 18 micron fiber length cellulose polymer coated with 0.25% by weight of the silicon containing compound simethicone;
- (ii) 500 ppm of the silicon containing compound simethicone in admixture with the polymer in (i);
- (iii) 100 ppm of the anti-gas agent lauryl sulfate;

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(iv) 0.3% by weight xanthan gum; and
(v) optionally, the flavoring agent Vivonex™
(commercially available from Norwhich Eaton Pharmaceutical
Inc., Norwhich, New York).

5 The present invention is further described in the
following Examples. In the Examples which follow, the
polymers described are in the form of solid fibers, with
any sizes given denoting fiber length. The Examples are
not to be construed with as limiting the scope of the
10 appended Claims.

EXAMPLES

Example 1

Surface tension, foaming and bubble formation was measured for various contrast media comprising
15 biocompatible polymers, both (i) with coating and/or in admixture with a silicon containing compound, and (ii) without coating and/or in admixture with a silicon containing compound. In some cases anti-gas agents and/or suspending agents were included in the contrast media
20 tested. All samples were freshly degassed prior to use by sonication using a commercially available sonicator (Bransen, 2200 Danbury, CT) under vacuum using a commercially available vacuum pump (Cole Parmer Model No. 7049-50, Chicago, IL).

25 Surface tension measurements were performed at 25°C using a CSC-DuNoüy tensiometer No. 70535 (Fairfax, VA). Degree of foaming was assessed by shaking the freshly degassed solutions in 50 cc plastic tubes (Fisher Scientific 05-539-8, Pittsburgh, PA). A volume of 25 cc of
30 each contrast agent suspension was placed in the tube and the tubes were capped. The tubes were then shaken vigorously for 60 seconds and the degree of foaming was assessed at a time-point (60 seconds) after the shaking ceased. Foam was recorded on the scale of 0 = no foam, 1 = minimal foam (about 1 mm), 2 = moderate foam (about 2 to 3 mm), 3 = severe foaming (more than 4 mm). The bubbles in the suspensions were assessed similarly after shaking but

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the bubbles were counted only within the suspensions, not as the foam which was determined by assessing the layer at the top of the suspension. Bubbles within the suspensions were assessed by examining the bubbles and grading them as 5 0 = no bubbles, 1 = small bubbles (<1 mm diameter with very few bubbles), 2 = medium bubbles (1-2 mm diameter and few to moderate bubbles), and 3 = large bubbles (>2 mm diameter with many bubbles).

10 The data from the surface tension, foaming, and bubble measurements are set forth in the Tables I-III. As shown in Table I, cellulose functions as a mild surfactant (that is, its surface tension is less than water), when the cellulose is prepared alone. When xanthan gum is added to the cellulose, this causes the mixture to lose some of its 15 favorable surfactant properties and the surface tension is similar to water. Polyethylene glycol (PEG 3350) has low surface tension and no foaming (but note in Example 2, Table IV, that PEG has no appreciable reflectivity on ultrasonic study). Methylcellulose caused appreciable 20 foaming. Simethicone coated cellulose, on the other hand, has low surface tension, no appreciable foaming, and no appreciable bubbles.

25 **Table I**
Effect of Simethicone/Silicone Coatings/Admixtures
on Surface Tension, Foaming and Bubble Formation

		Dynes/cm	Foam	Bubbles
	Deionized Water	73.5	0.0	0.0
	18 μ Cellulose			
30	1% cellulose	71.3	0.0	0.0
	2% cellulose	71.2	0.0	0.0
	18 μ cellulose w/0.25% xanthan gum			
	1% cellulose	72.7	0.0	1.0
	2% cellulose	72.8	0.0	1.0

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Table I (cont'd)

		Dynes/cm	Foam	Bubbles
	18μ cellulose w/0.05% xanthan gum			
5	1% cellulose	71.6	0.0	0.5
	2% cellulose	72.0	0.0	0.05
	18μ cellulose 1% simethicone coated 0.25% xanthan gum	70.6	0.0	0.0
10	18μ cellulose 1% silicone coated 0.25% xanthan gum	70.3	0.0	0.0
	1% cellulose 1% silicone coated	61.0	0.0	0.0
15	2% cellulose 1% silicone coated	61.8	0.0	0.0
	1% cellulose 1% simethicone coated	62.0	0.0	0.0
20	2% cellulose 1% simethicone coated	61.0	0.0	0.0

Key:

Foam 0.0 - 3.0
 Bubbles 0.0 - 3.0
 (for foam and bubbles,
 zero is the most favorable)
 Dynes (20.0 - 80.0)
 (for dynes, the lower the number,
 the better)

Table II

30 Effect of Anti-Gas Agents
 on Surface Tension, Foaming and Bubble Formation
 of Simethicone/Silicone Coatings/Admixtures

		Dynes/cm	Foam	Bubbles
	sodium lauryl sulfate w/ 1% 18μ 1% silicone coated cellulose w/ 0.15% xanthan gum			
35	100 ppm sodium lauryl sulfate	49.1	1.0	1.0
40				

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Table II (cont'd)

	Dynes/cm	Foam	Bubbles
--	----------	------	---------

5	500 ppm sodium lauryl sulfate	34.4	2.0	1.5
10	Sodium lauryl sulfate w/ 1% 18μ 1% silicone coated cellulose w/ 0.15% xanthan gum			
15	100 ppm sodium lauryl sulfate	52.2	1.0	1.0
20	500 ppm sodium lauryl sulfate	35.1	2.0	1.5
25	1000 ppm Simethicone w/500 ppm sodium lauryl sulfate w/ 1% 18μ 1% silicone coated cellulose w/ 0.15% xanthan gum	32.9	0.0	0.0
30	Key:	Foam 0.0 - 3.0 Bubbles 0.0 - 3.0 (for foam and bubbles, zero is the most favorable) Dynes (20.0 - 80.0) (for dynes, the lower the number, the better)		
35				

Table III

Comparison of Various Polymer Compositions
in Surface Tension, Foaming and Bubble Formation

40		Dynes/cm	Foam	Bubbles
	Deionized Water	73.5	0.0	0.0
	18μ Cellulose 1% cellulose	71.3	0.0	0.0

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Table III (cont'd)

		Dynes/cm	Foam	Bubbles
	2% cellulose	71.2	0.0	0.0
5	Polyethylene Glycol (PEG3350). (Not Degassed)			
	1% PEG	63.3	0.0	0.0
	2% PEG	62.8	0.0	0.0
	3% PEG	63.1	0.0	0.0
	4% PEG	62.8	0.0	0.0
10	5% PEG	62.4	0.0	0.0
	Polyethylene Glycol (PEG3350) (Degassed)			
	1% PEG	62.3	0.0	0.0
	2% PEG	62.1	0.0	0.0
15	3% PEG	61.6	0.0	0.0
	4% PEG	61.5	0.0	0.0
	5% PEG	61.2	0.0	0.0
	18μ Cellulose			
20	w/ 0.25% Xanthan Gum			
	1% cellulose	72.7	0.0	1.0
	2% cellulose	72.8	0.0	1.0
	3% cellulose	73.1	0.0	1.0
	4% cellulose	72.8	0.0	1.0
	5% cellulose	73.0	0.0	1.0
25	18μ Cellulose			
	w/ 0.05% Xanthan Gum			
	1% cellulose	71.6	0.0	0.5
	2% cellulose	72.0	0.0	0.5
	3% cellulose	71.7	0.0	0.5
30	4% cellulose	71.8	0.0	0.5
	5% cellulose	71.9	0.0	0.5
	Polyethylene Glycol (PEG3350)			
35	w/ 1% 18μ Cellulose			
	w/ 0.15% Xanthan Gum			
	1% PEG	54.1	0.0	1.5
	2% PEG	51.2	0.0	1.5
	3% PEG	56.0	0.0	1.5
40	18μ Cellulose			
	1% Simethicone Coated			
	w/ 0.25% Xanthan Gum	70.6	0.0	0.0
	18μ Cellulose			
	2% Silicone Coated			
	0.25% Xanthan Gum	70.3	0.0	0.0

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Table III (cont'd)

		Dynes/cm	Foam	Bubbles
5	Propylene Glycol w/ 1% 18μ 1% Silicone Coated Cellulose w/ 0.15% Xanthan Gum 3% PG 5% PG	61.8 62.0	0.0 0.0	1.0 1.5
10	Sodium Lauryl Sulfate w/ 1% 18μ 1% Silicone Coated Cellulose w/ 0.15% Xanthan Gum 100 ppm SLS 500 ppm SLS	49.1 34.4	1.0 2.0	1.0 1.5
15	Sodium Lauryl Sulfate w/ 2% Polyethylene Glycol (PEG3350) w/ 1% 18μ 1% Silicone Coated Cellulose			
20	w/ 0.15% Xanthan Gum 100 ppm SLS 500 ppm SLS	51.5 34.5	1.0 2.0	1.5 2.0
25	Sodium Lauryl Sulfate w/ 1% 18μ 1% Silicone Coated Cellulose w/ 0.15% Xanthan Gum 100 ppm SLS 500 ppm SLS	52.2 35.1	1.0 2.0	1.0 1.5
30	Simethicone w/ 500 ppm Sodium Lauryl Sulfate w/ 1% 18μ Silicone Coated Cellulose w/ 0.15% Xanthan Gum 1000 ppm simethicone	32.9	0.0	1.0
35	1% 18μ 1% Simethicone Coated Cellulose w/ 500 ppm			
40	Sodium Lauryl Sulfate w/ 1000 ppm Simethicone w/ 0.15% Xanthan Gum #1 #2 #3	30.8 33.1 33.2	1.5 1.0 1.0	1.5 1.5 1.5
45	#4 #5 #6 #7 #8	33.2 32.9 33.2 32.8 33.1	0.5 0.5 0.5 0.5 0.5	1.5 1.5 1.5 1.5 1.5

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Table III (cont'd)

		Dynes/cm	Foam	Bubbles
	#9	32.1	0.5	1.5
	Methyl Cellulose			
5	0.15%	54.6	2.0	0.0
	0.25%	53.9	2.5	0.0
	0.50%	54.1	2.5	0.0
	1.00%	54.5	3.0	0.0
	Hydroxyethyl Cellulose (Natrasol™)			
10	0.15%	64.0	2.0	0.0
	0.25%	56.1	1.5	0.5
	0.50%	51.7	1.0	1.5
	1.00%	50.4	0.5	2.5
	Carrageenan			
15	0.15%	72.9	0.0	0.0
	0.25%	72.6	0.0	0.0
	0.50%	72.8	0.0	0.5
	Xanthan Gum			
20	0.15%	72.8	0.0	0.5
	0.25%	73.6	0.0	1.0
	0.50%	73.1	0.0	1.0
	1.00%	75.9	0.0	1.5
	Carboxymethyl Cellulose			
25	0.15%	72.0	0.0	0.5
	0.25%	72.4	0.0	1.0
	0.50%	72.4	0.0	1.5
	1.00%	73.3	0.0	2.0
	Sodium Lauryl Sulfate			
30	100 ppm	51.4	1.0	0.0
	200 ppm	48.6	1.5	0.0
	300 ppm	35.7	2.0	0.0
	400 ppm	29.8	2.5	0.0
	500 ppm	29.5	3.0	0.0
	Corn Starch			
35	w/ 0.15% Xanthan Gum			
	1% corn starch	73.0	0.0	0.0
	3% corn starch	73.1	0.0	0.0
	5% corn starch	73.1	0.0	0.0
	Wheat (Cook-up) Starch			
40	w/ 0.15% Xanthan Gum			
	1% wheat starch	72.2	0.0	1.5
	3% wheat starch	71.0	0.0	2.0
	5% wheat starch	69.7	0.0	2.5

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Table III (cont'd)

		Dynes/cm	Foam	Bubbles
	Wheat (Instant) Starch w/ 0.15% Xanthan Gum			
5	1% wheat starch	59.9	0.0	2.0
	3% wheat starch	56.7	0.0	2.0
	5% wheat starch	----	0.0	2.5
	1% 18 μ 1% Simethicone			
10	Coated Cellulose w/ 0.30% Xanthan Gum (Keltrol RD TM)			
	w/ 100 ppm			
	Sodium Lauryl Sulfate			
15	w/ 500 ppm Simethicone			
	w/ 0.5% flavoring	37.6	0.0	0.0
	Key:	Foam 0.0 - 3.0 Bubbles 0.0 - 3.0 (for foam and bubbles, zero is the most favorable) Dynes (20.0 - 80.0; (for dynes, the lower the number,		
20				

Example 2

- Velocity, acoustic impedance, dB reflectivity, and attenuation was measured for various contrast media comprising biocompatible polymers, both (i) with coating and/or in admixture with a silicon containing compound, and (ii) without coating and/or in admixture with a silicon containing compound.
- The dB reflectivity of the different contrast agents was assessed with an Acoustic Imaging Model 5200 ultrasound scanner (Phoenix, AZ) equipped with Performance Software PFM. The transducers, either 5.0 or 7.5 Mhz, were placed in the contrast agents and scanned. Prior to performing each set of experiments the dB reflectivity was standardized to a known reference phantom. For the dB reflectivity measurements on the Acoustic Imaging 5200, a tissue mimicking phantom was used as a reference. The tissue mimicking phantom is made by Nuclear Associates. The gain was set to -53.5 dB. The dB reflectivity measurements were made by selecting a region of interest on

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the CRT monitor and reading the mean dB reflectivity in that region. Quantitative measurements of samples were obtained on a benchtop acoustic lab custom built by ImaRx, (Tucson, AZ), consisting of a Panametric (Waltham, MA) 5 5052PR ultrasonic pulser/receiver, LeCroy (Chestnut Ridge, NY) 9410 dual channel digital oscilloscope with a waveform processing package and capabilities of Fast Fourier Transform and Krautkramer Branson (Lewistown, PA), F-style, Gamma-HP series transducers. Data was obtained using the 10 pulse-echo technique. The returned echoes were measured with the appropriate time delays and the power average of the waveform was calculated. The frequencies of the transducers used were 2.25, 3.25, 5.0, 7.5 and 10 mHz. The 15 ultrasonic waves were reflected from a solution/air interface reflected from a solution/air interface. Parameters of velocity, acoustic impedance, attenuation and transmittance were assessed.

The data from the benchtop acoustic lab are set forth in Table IV. As shown by these data, cellulose with 20 simethicone coating has both good reflectivity and very high transmission (that is, less acoustic attenuation) than plain cellulose. Note also that polyethyleneglycol has much lower reflectivity than cellulose.

Figures 4-8 show the dB reflectivity from the 25 measurements made on the phantoms with the Acoustic Imaging Ultrasound machine. As shown by these figures, for plain cellulose the shorter fibers (e.g. 18 through 35 micron) have higher reflectivity than the longer cellulose fibers. Note that the silicone coated cellulose has higher 30 reflectivity than the uncoated fibers.

Example 3

Hexamethyldisilazane (1,1,1,3,3,3-hexamethyldisilazane) (2 g), commercially available from Aldrich Chemical Company, Milwaukee Wisconsin, was added to 35 10g of ethyl alcohol (200 proof dehydrated alcohol, U.S.P. punctilious). The hexamethyldisilazane and ethyl alcohol mixture was then added drop by drop to 28.5 g of cellulose

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(fiber lengths 18 μ , 22 μ , and 35 μ , respectively) in a flask. The mixture was stirred and heated at a temperature of about 60°C in a vacuum oven. After the ethyl alcohol was evaporated, the flask was placed into a laboratory oven and 5 heated from about 60°C to about 140°C for about two hours and then cooled, resulting in hexamethyldisilazane coated cellulose (also referred to herein generally as silylated cellulose).

Example 4

10 Silicone (in oil form) (2 g), commercially available from Aldrich Chemical Co. Inc., was added to 10 g of toluene. The silicone and toluene mixture was then added drop by drop to 28.5 g of cellulose in a flask. The mixture was stirred and heated at a temperature of about 15 60°C in a vacuum oven. After the toluene was evaporated, the flask was placed into a laboratory oven and heated from about 60°C to about 140°C for about two hours and then cooled, resulting in silicone coated cellulose.

Example 5

20 A solution containing about 15% by weight polyethylene glycol fibers having a molecular weight of about 8000 was prepared in deionized water. The solution was then mixed with a gas and the solution was scanned in vitro by ultrasound.

25 The polyethylene glycol polymer solution was found to improve gas dissipation and render a good image on in vitro scanning.

Example 6

30 Three solutions containing about 5% by weight cellulose fibers having varying fiber lengths of cellulose of about 22, 60, and 110 microns, respectively, and each containing about 0.25% xanthan gum were prepared in distilled water. A fourth solution containing about 5% by weight dextran beads of about 20 micron diameter was 35 prepared in distilled water. The solutions were scanned in vitro by ultrasound at constant gain settings using a 5

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megahertz transducer (approximately a 300 micron wave length in aqueous media).

The 20 micron diameter dextran bead solution had the highest echogenicity of the four solutions, with the 22 5 micron fiber length cellulose solution having the next highest echogenicity.

Example 7

A solution containing about 5% by weight cellulose fibers having a fiber length of about 22 microns 10 and about 0.25% by weight xanthan gum was prepared in deionized water. Approximately 750 ml of the solution was then administered orally to a mammal, and ultrasonic imaging of the gastrointestinal tract was performed.

Distention of the gastrointestinal tract and good 15 visualization of the gastrointestinal region was achieved. Excellent visualization of the mucosal surfaces of the stomach and intestine was also achieved.

Example 8

A solution containing about 4% by weight 20 cellulose fibers having a fiber length of about 22 microns and about 2 g of activated charcoal in deionized water was prepared. Approximately 750 ml of the solution was then administered orally to a mammal and ultrasonic imaging of the gastrointestinal tract was performed.

25 Ultrasonic imaging provided good visualization of the tract and mucosal surfaces.

Example 9

A solution containing about 5% by weight cellulose fibers having a fiber length of about 22 microns 30 and about 0.25% by weight xanthan gum was prepared in deionized water. To this solution, about 2.0 g of simethicone was added and the solution was mixed. Approximately 750 ml of the solution was then administered orally to a mammal, and ultrasonic imaging of the gastro- 35 intestinal tract was performed.

Ultrasonic imaging provided good visualization of the tract and mucosal surfaces. In addition, improved

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removal of gas bubbles from the gastrointestinal tract was observed.

Various modifications of the invention in addition to those shown and described herein will be apparent to those skilled in the art from the foregoing description. Such modifications are also intended to fall within the scope of the appended Claims.

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CLAIMS

What is claimed is:

1. A contrast medium for ultrasonic imaging comprising an aqueous solution of at least one biocompatible polymer, wherein the biocompatible polymer is coated with and/or in admixture with at least one silicon containing compound.
2. A contrast medium of Claim 1 wherein the polymer is a synthetic polymer.
3. A contrast medium of Claim 1 wherein the polymer is a natural polymer or a semisynthetic polymer.
4. A contrast medium of Claim 2 wherein the polymer is selected from the group consisting of polyethylenes, polypropylenes, pluronic acids, pluronic alcohols, polyvinyls, nylon, polystyrene, polylactic acids, fluorinated hydrocarbons, fluorinated carbons, polydimethylsiloxane, and polymethylmethacrylate.
5. A contrast medium of Claim 3 wherein the polymer is selected from the group consisting of arabinans, fructans, fucans, galactans, galacturonans, glucans, mannans, xylans, levan, fucoidan, carrageenan, galactocarolose, pectic acid, amylose, pullulan, glycogen, amylopectin, cellulose, carboxymethylcellulose, hydroxymethyl cellulose, hydroxypropyl methylcellulose, methyl cellulose, methoxycellulose, dextran, pustulan, chitin, agarose, keratan, chondroitin, dermatan, hyaluronic acid, alginic acid, xanthan gum, and polysaccharides containing at least one aldose, ketose, acid or amine selected from the group consisting of erythrose, threose, ribose, arabinose, xylose, lyxose, allose, altrose, glucose, mannose, gulose, idose, galactose, talose, erythrulose, ribulose, xylulose, psicose, fructose, sorbose, tagatose, glucuronic acid, mannuronic acid, glucosamine, galactosamine, and neuraminic acid.
6. A contrast medium of Claim 1 wherein the polymer has a high water binding capacity.

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7. A contrast medium of Claim 1 further comprising an anti-gas agent.

8. A contrast medium of Claim 7 wherein the anti-gas agent is selected from the group consisting of activated charcoal, aluminum carbonate, aluminum hydroxide, aluminum phosphate, calcium carbonate, dihydroxyaluminum sodium carbonate, magaldrate, magnesium oxide, magnesium trisilicate, sodium carbonate, loperamide hydrochloride, diphenoxylate, hydrochloride with atropine sulfate, kaolin, bismuth salts, polyoxypropylene-polyoxyethylene copolymers, polyoxyalkylene amines and imines, branched polyamines, mixed oxyalkylated alcohols, sucroglycamides, polyoxylalkylated natural oils, lauryl sulfates, 2-lactylic acid esters of unicarboxylic acids, triglyceride oils, and finely-divided polyvinyl chloride particles.

9. A contrast medium of Claim 1 further comprising a suspending agent.

10. A contrast medium of Claim 9 wherein the suspending agent is selected from the group consisting of xanthum gum, acacia, agar, alginic acid, aluminum monostearate, unpurified bentonite, purified bentonite, bentonite magma, carbomer 934P, carboxymethylcellulose calcium, carboxymethylcellulose sodium, carboxymethyl-cellulose sodium 12, carrageenan, cellulose (microcrystalline), carboxymethylcellulose sodium, dextrin, gelatin, guar gum, hydroxyethyl cellulose, hydroxypropyl cellulose, hydroxypropyl methylcellulose, magnesium aluminum silicate, methylcellulose, pectin, polyethylene oxide, polyvinyl alcohol, povidone, propylene glycol alginate, silicon dioxide, silicon dioxide colloidal, sodium alginate, and tragacanth.

11. A contrast medium of Claim 1 wherein the silicon containing compound is selected from the group consisting of silicone, simethicone, protected simethicone, polysiloxane, polydimethylsiloxane, polymethylvinyl-siloxane, polymethylphenolsiloxane, polydiphenylsiloxane, octamethylcyclotetrasiloxane, siloxane glycol polymer,

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gamma-aminopropyltriethoxysilane, diethylaminomethyltriethoxysilane, silicon dioxide, siloxyalkylene polymer, hexadimethylsilazane, hexamethyldisilazane, hexaphenylcyclotrisilazane, octamethylcyclotetrasilazane, N'N'-bis(trimethylsilyl)acetamide, N-(dimethyl(gamma-cyanopropyl)-silyl)-N-methylacetamide), and siliceous earth.

12. A contrast medium of Claim 1 wherein the polymer is coated with at least one silicon containing compound.

13. A contrast medium of Claim 1 wherein the aqueous solution is degassed and stored under vacuum.

14. A contrast medium of Claim 1

wherein the polymer comprises 0.5% by weight of an 18 μ fiber length cellulose polymer; and

wherein the cellulose polymer is coated with 0.25% by weight of the silicon containing compound simethicone and admixed with 500 ppm of the silicon containing compound simethicone; and

further comprising an anti-gas agent comprising lauryl sulfate and a suspending agent comprising 0.3% by weight xanthan gum; and

wherein the aqueous solution is degassed and the contrast medium is stored under vacuum.

15. A contrast medium comprising an aqueous solution of at least one biocompatible synthetic polymer.

16. A contrast medium of Claim 15 wherein the polymer is selected from the group consisting of polyethylenes, polypropylenes, pluronic acids, pluronic alcohols, polyvinyls, nylon, polystyrene, polylactic acids, fluorinated hydrocarbons, fluorinated carbons, polydimethylsiloxane, and polymethylmethacrylate.

17. A contrast medium of Claim 15 wherein the polymer has a high water binding capacity.

18. A contrast medium of Claim 15 further comprising an anti-gas agent.

19. A contrast medium of Claim 18 wherein the anti-gas agent is selected from the group consisting of

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activated charcoal, aluminum carbonate, aluminum hydroxide, aluminum phosphate, calcium carbonate, dihydroxyaluminum sodium carbonate, magaldrate, magnesium oxide, magnesium trisilicate, sodium carbonate, loperamide hydrochloride, diphenoxylate, hydrochloride with atropine sulfate, kaolin, bismuth salts, polyoxypropylene-polyoxyethylene copolymers, polyoxyalkylene amines and imines, branched polyamines, mixed oxyalkylated alcohols, sucroglycamides, polyoxylalkylated natural oils, lauryl sulfates, 2-lactylic acid esters of unicarboxylic acids, triglyceride oils, and finely-divided polyvinyl chloride particles.

20. A contrast medium of Claim 15 further comprising a suspending agent.

21. A contrast medium of Claim 20 wherein the suspending agent is selected from the group consisting of xanthan gum, acacia, agar, alginic acid, aluminum monostearate, unpurified bentonite, purified bentonite, bentonite magma, carbomer 934P, carboxymethylcellulose calcium, carboxymethylcellulose sodium, carboxymethylcellulose sodium 12, carrageenan, cellulose (microcrystalline), carboxymethylcellulose sodium, dextrin, gelatin, guar gum, hydroxyethyl cellulose, hydroxypropyl cellulose, hydroxypropyl methylcellulose, magnesium aluminum silicate, methylcellulose, pectin, polyethylene oxide, polyvinyl alcohol, povidone, propylene glycol alginate, silicon dioxide, silicon dioxide colloidal, sodium alginate, and tragacanth.

22. A contrast medium of Claim 15 wherein the aqueous solution is degassed and stored under vacuum.

23. A contrast medium comprising an aqueous solution of cellulose.

24. A contrast medium of Claim 23 further comprising an anti-gas agent.

25. A contrast medium of Claim 24 wherein the anti-gas agent is selected from the group consisting of activated charcoal, aluminum carbonate, aluminum hydroxide, aluminum phosphate, calcium carbonate, dihydroxyaluminum

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- sodium carbonate, magaldrate, magnesium oxide, magnesium trisilicate, sodium carbonate, loperamide hydrochloride, diphenoxylate, hydrochloride with atropine sulfate, kaolin, bismuth salts, polyoxypropylene-polyoxyethylene copolymers, polyoxyalkylene amines and imines, branched polyamines, mixed oxyalkylated alcohols, sucroglycamides, polyoxylalkylated natural oils, lauryl sulfates, 2-lactylic acid esters of unicarboxylic acids, triglyceride oils, and finely-divided polyvinyl chloride particles.

26. A contrast medium of Claim 23 further comprising a suspending agent.

27. A contrast medium of Claim 26 wherein the suspending agent is selected from the group consisting of xanthan gum, acacia, agar, alginic acid, aluminum monostearate, unpurified bentonite, purified bentonite, bentonite magma, carbomer 934P, carboxymethylcellulose calcium, carboxymethylcellulose sodium, carboxymethyl-cellulose sodium 12, carrageenan, cellulose (microcrystalline), carboxymethylcellulose sodium, dextrin, gelatin, guar gum, hydroxyethyl cellulose, hydroxypropyl cellulose, hydroxypropyl methylcellulose, magnesium aluminum silicate, methylcellulose, pectin, polyethylene oxide, polyvinyl alcohol, povidone, propylene glycol alginate, silicon dioxide, silicon dioxide colloidal, sodium alginate, and tragacanth.

28. A contrast medium of Claim 23 wherein the aqueous solution is degassed and stored under vacuum.

29. A method of providing an image of an internal region of a patient comprising (i) administering to the patient a contrast medium of Claim 1, and (ii) scanning the patient using ultrasonic imaging to obtain visible images of the region.

30. A method for diagnosing the presence of diseased tissue in a patient comprising (i) administering to the patient a contrast medium of Claim 1 and (ii) scanning the patient using ultrasonic imaging to obtain visible images of any diseased tissue in the patient.

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31. A method of providing an image of an internal region of a patient comprising (i) administering to the patient a contrast medium of Claim 15, and (ii) scanning the patient using ultrasonic imaging to obtain visible images of the region.

32. A method for diagnosing the presence of diseased tissue in a patient comprising (i) administering to the patient a contrast medium of Claim 15 and (ii) scanning the patient using ultrasonic imaging to obtain visible images of any diseased tissue in the patient.

33. A method of providing an image of an internal region of a patient comprising (i) administering to the patient a contrast medium of Claim 23, and (ii) scanning the patient using ultrasonic imaging to obtain visible images of the region.

34. A method for diagnosing the presence of diseased tissue in a patient comprising (i) administering to the patient a contrast medium of Claim 23 and (ii) scanning the patient using ultrasonic imaging to obtain visible images of any diseased tissue in the patient.

35. A kit for ultrasonic imaging comprising a contrast medium of Claim 1.

36. A kit in accordance with Claim 35 further comprising conventional ultrasonic imaging components wherein the conventional ultrasonic imaging components are selected from the group consisting of anti-gas agents, suspending agents, flavoring materials, coloring materials, antioxidants, anticaking agents, tonicity agents, and wetting agents.

37. A kit for ultrasonic imaging comprising a contrast medium of Claim 15.

38. A kit in accordance with Claim 37 further comprising conventional ultrasonic imaging components wherein the conventional ultrasonic imaging components are selected from the group consisting of anti-gas agents, suspending agents, flavoring materials, coloring materials,

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antioxidants, anticaking agents, tonicity agents, and wetting agents.

39. A kit for ultrasonic imaging comprising a contrast medium of Claim 23.

40. A kit in accordance with Claim 39 further comprising conventional ultrasonic imaging components wherein the conventional ultrasonic imaging components are selected from the group consisting of anti-gas agents, suspending agents, flavoring materials, coloring materials, antioxidants, anticaking agents, tonicity agents, and wetting agents.

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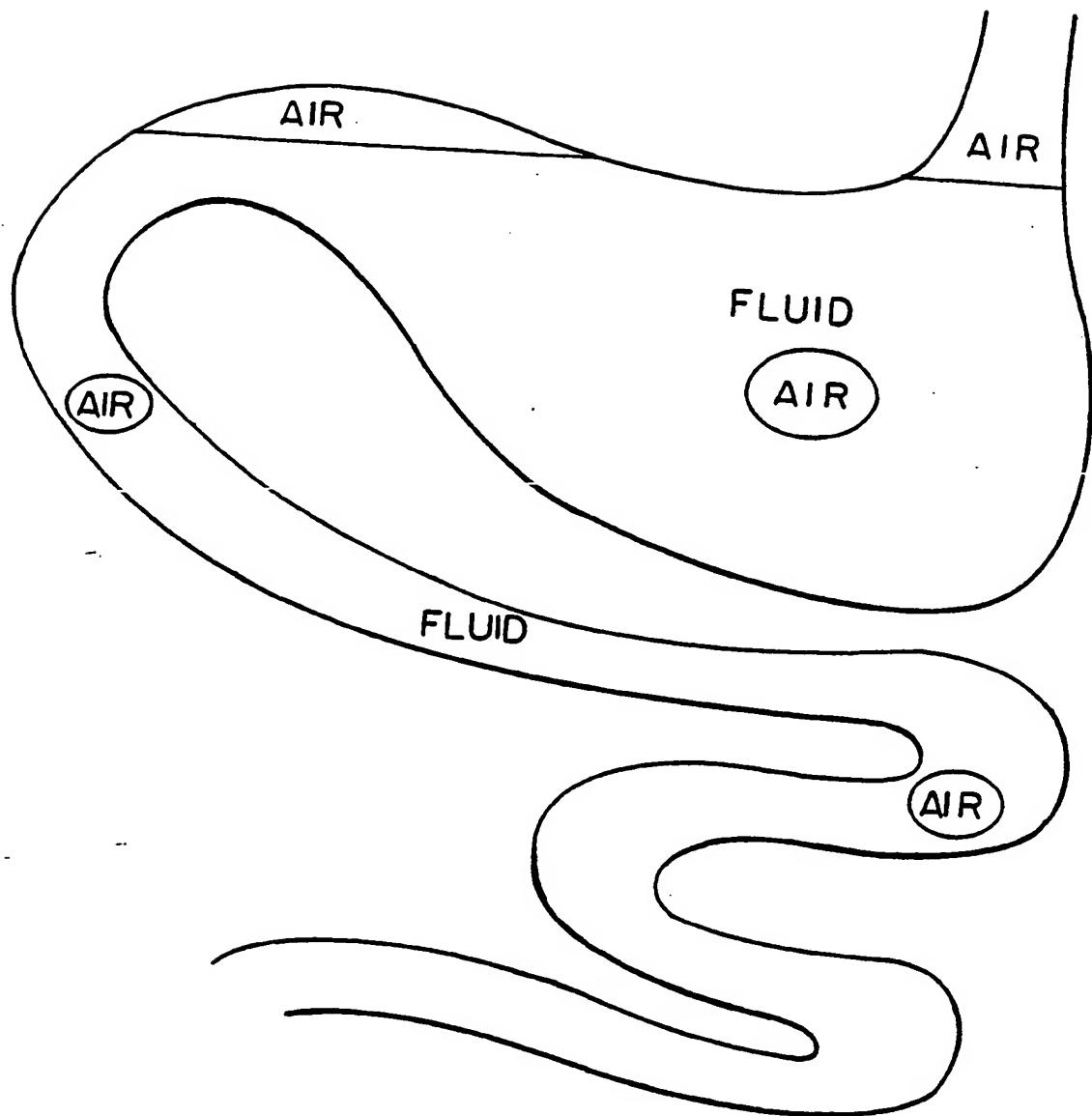


FIG. 1

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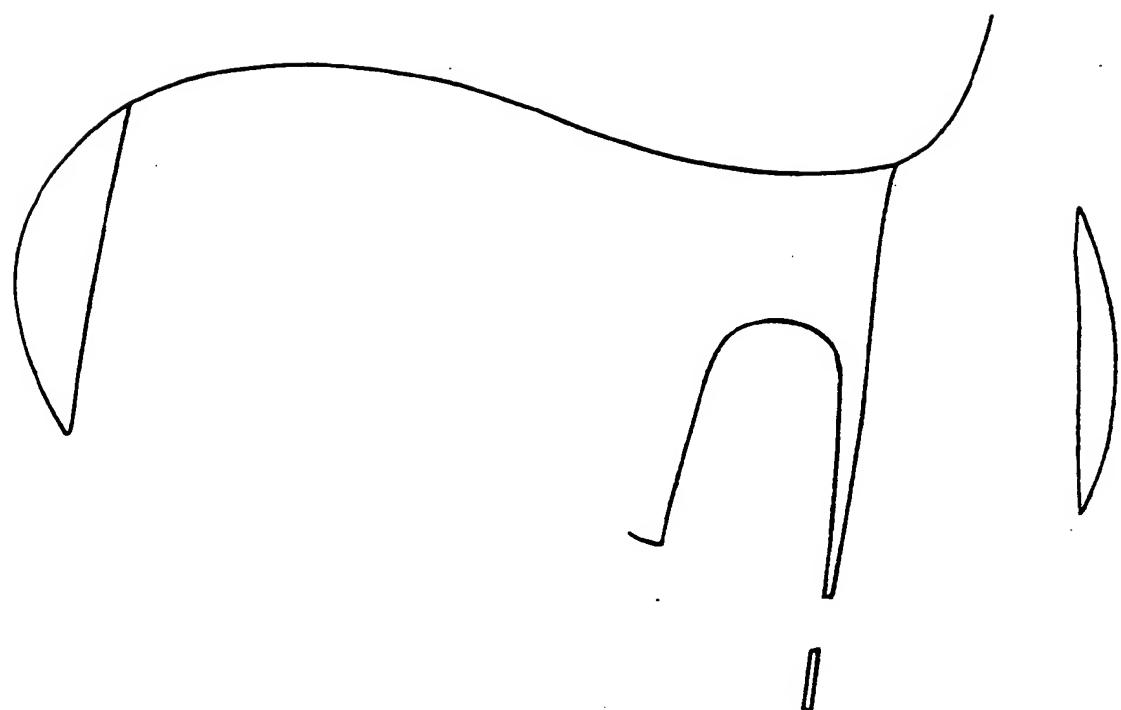


FIG. 2A

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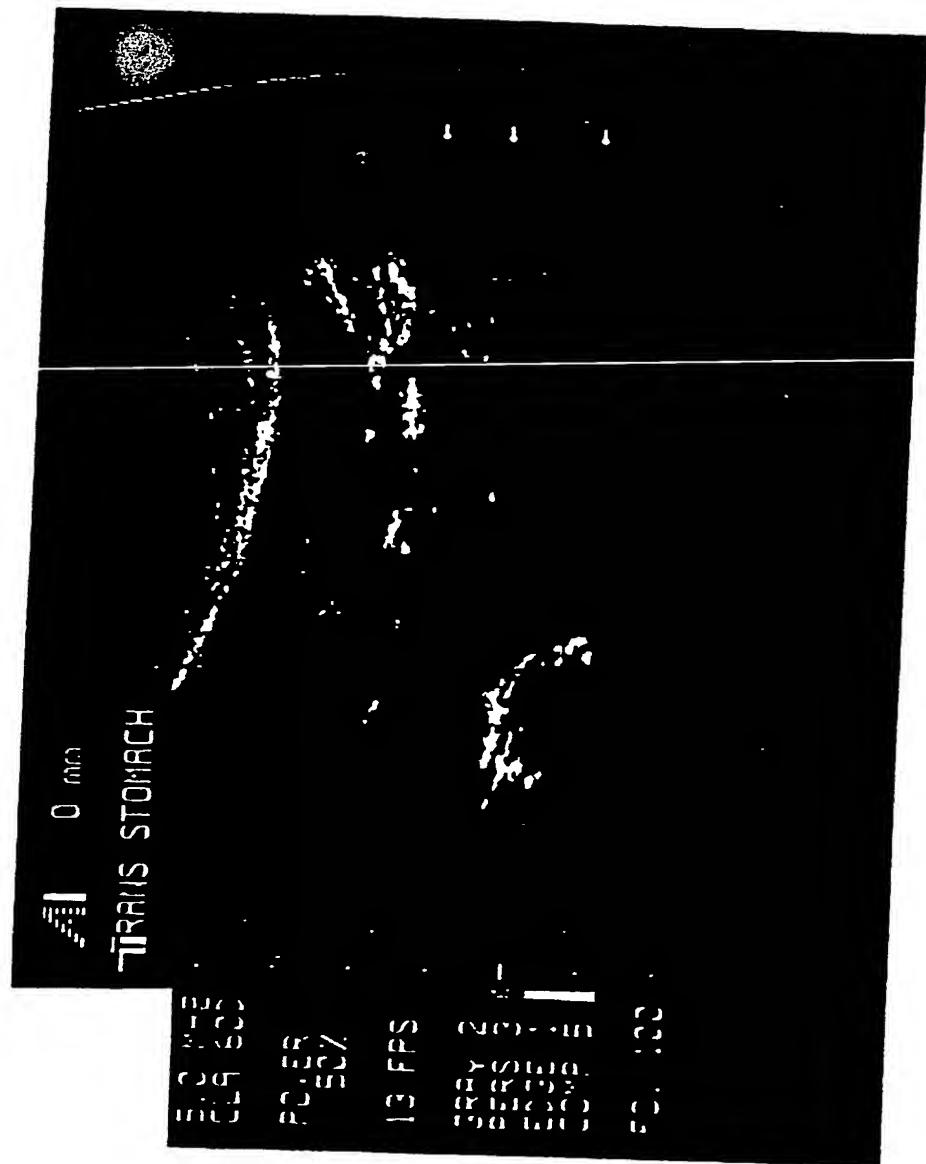


FIG. 2B

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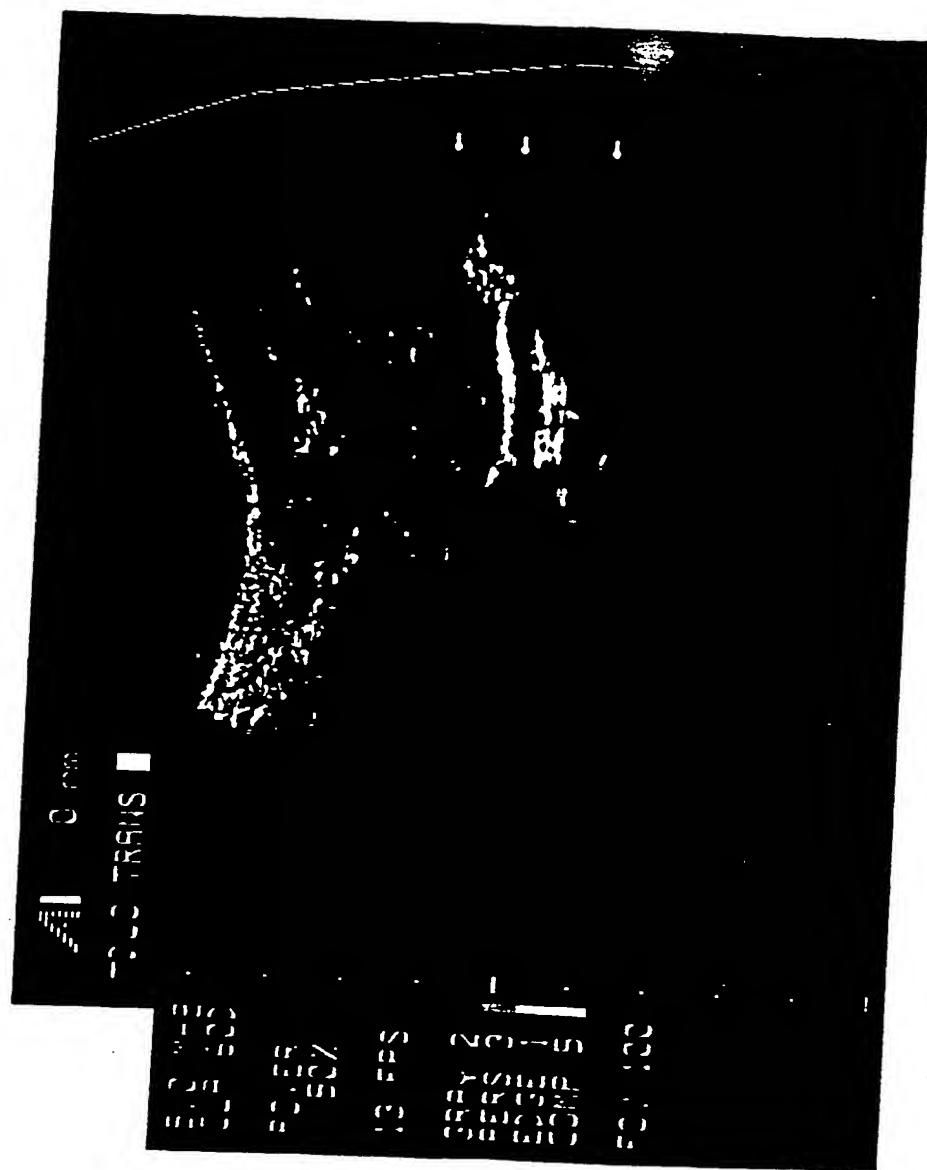


FIG. 2C

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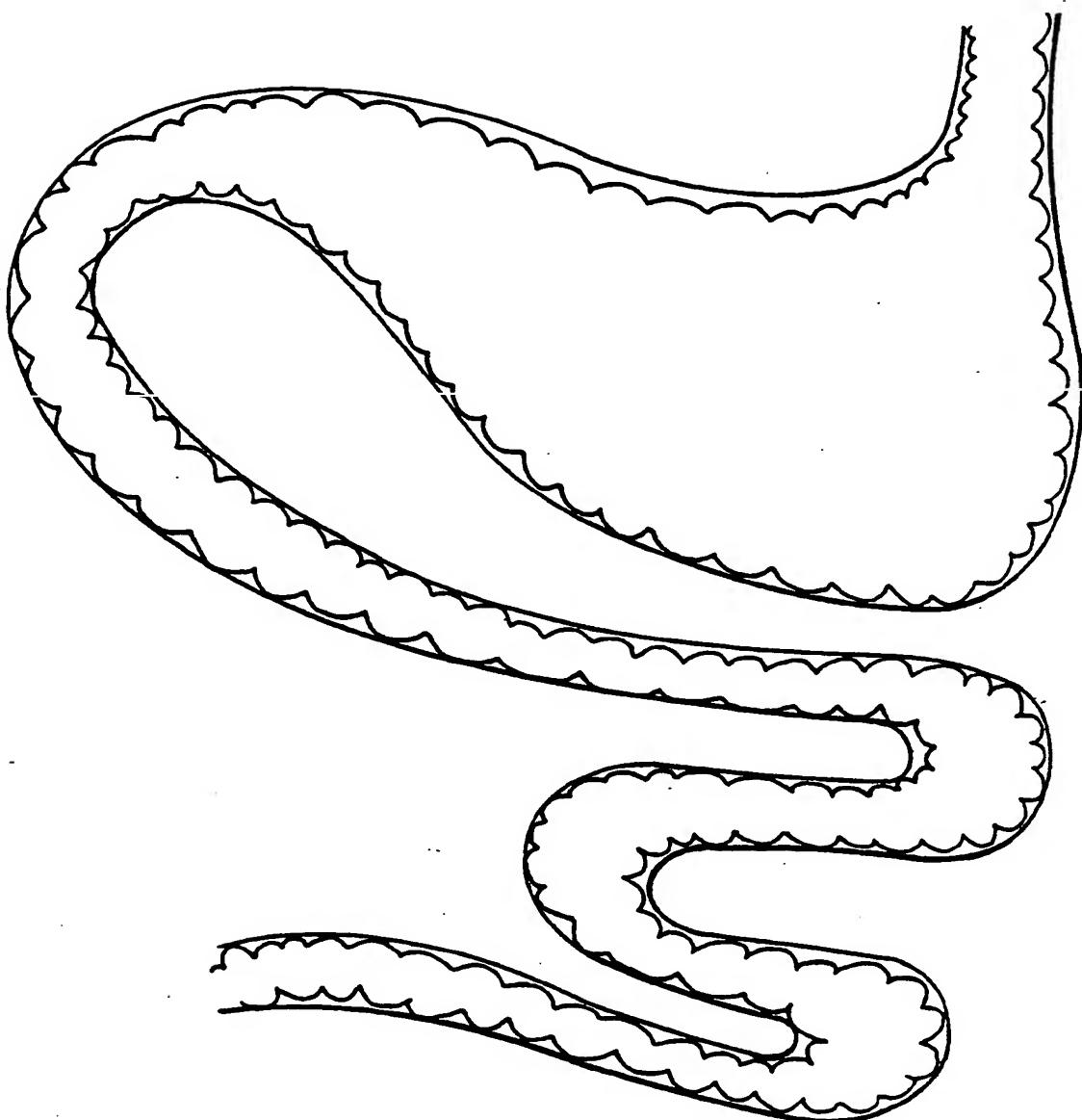


FIG. 3A

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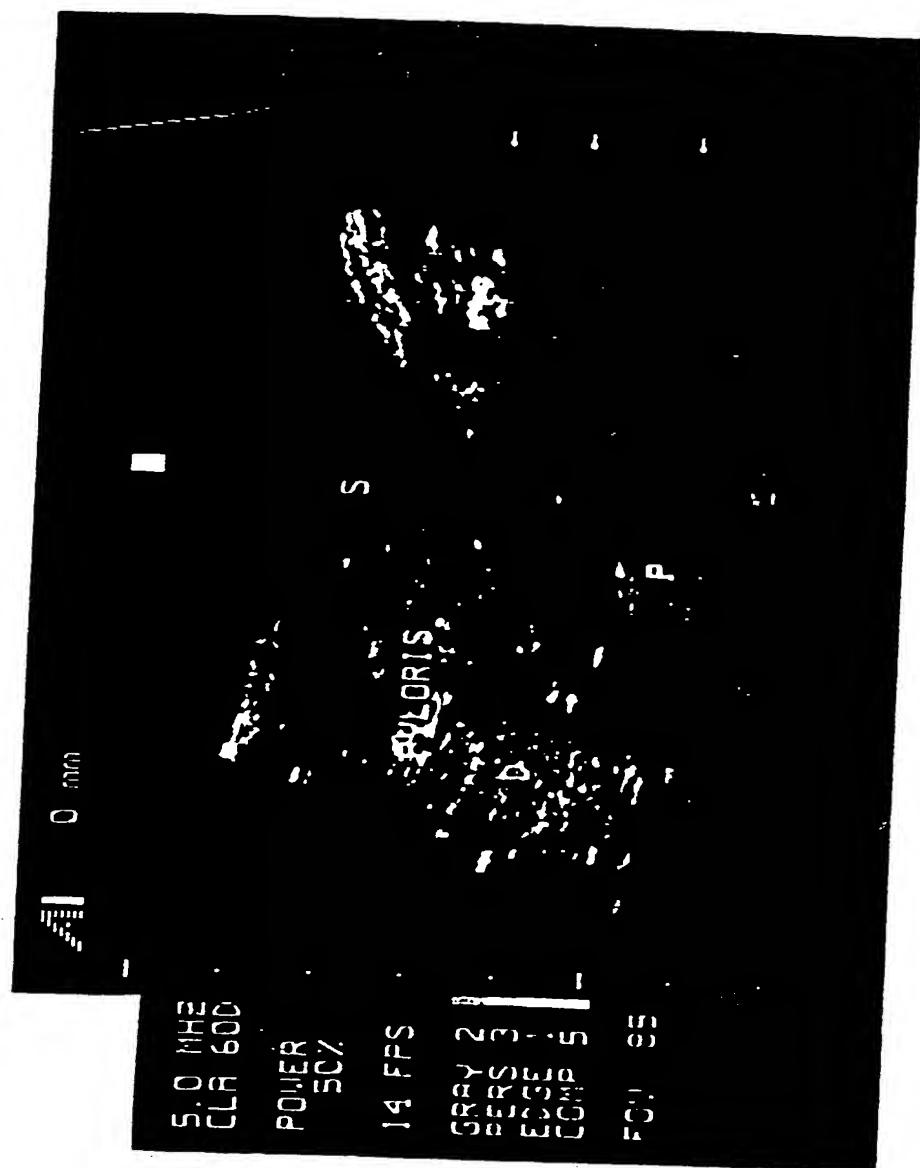


FIG. 3B

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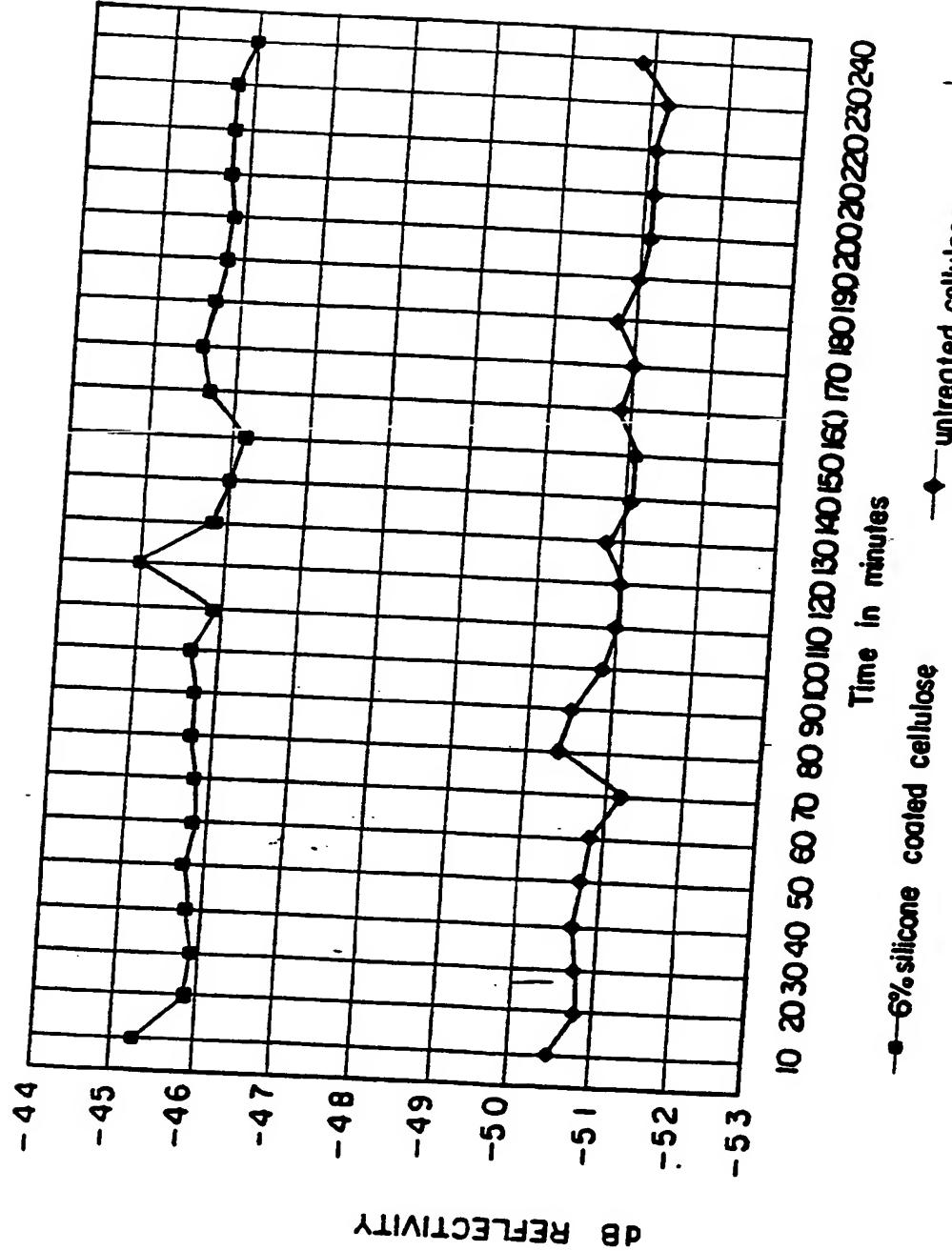


FIG. 4

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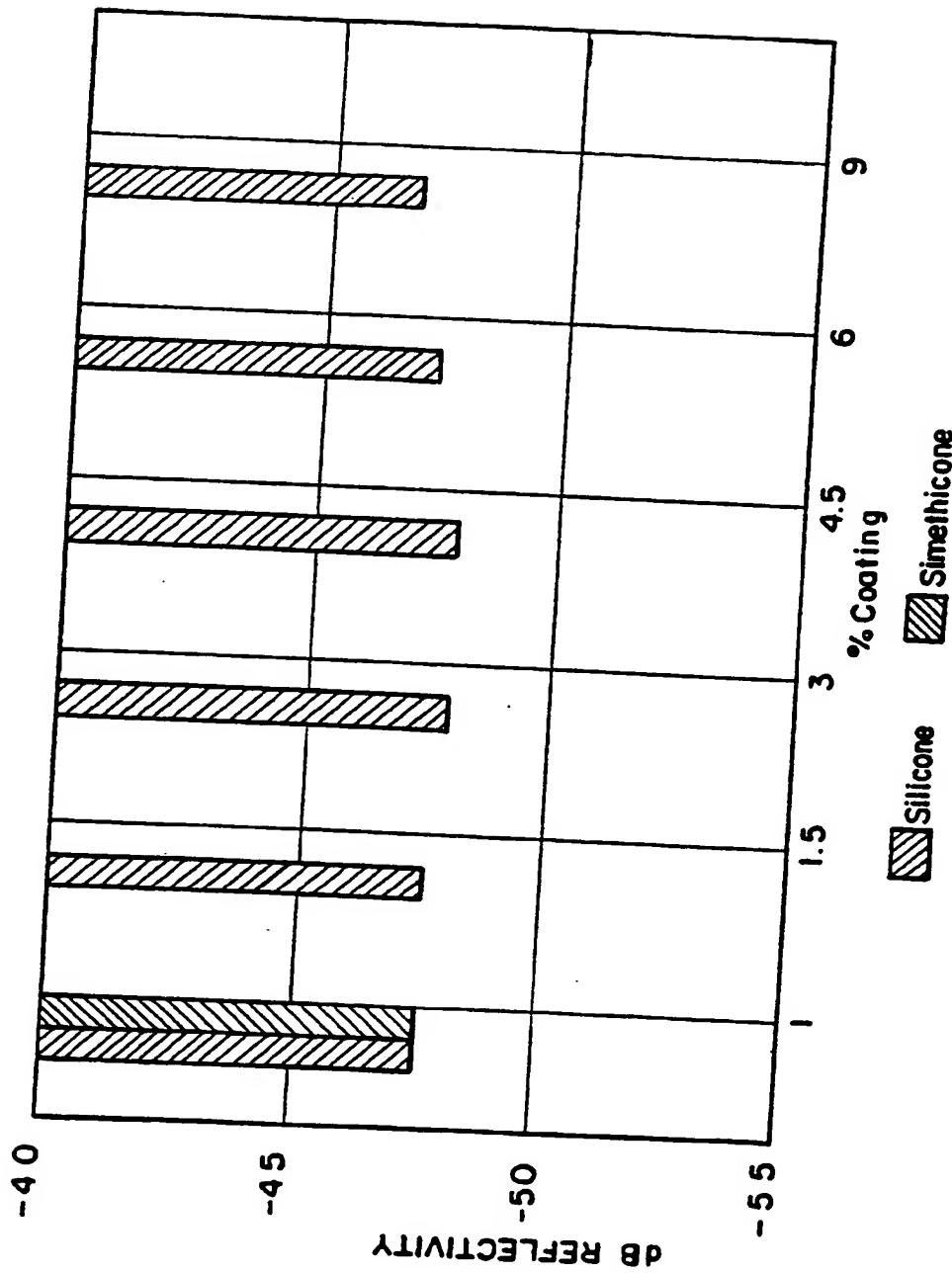


FIG. 5

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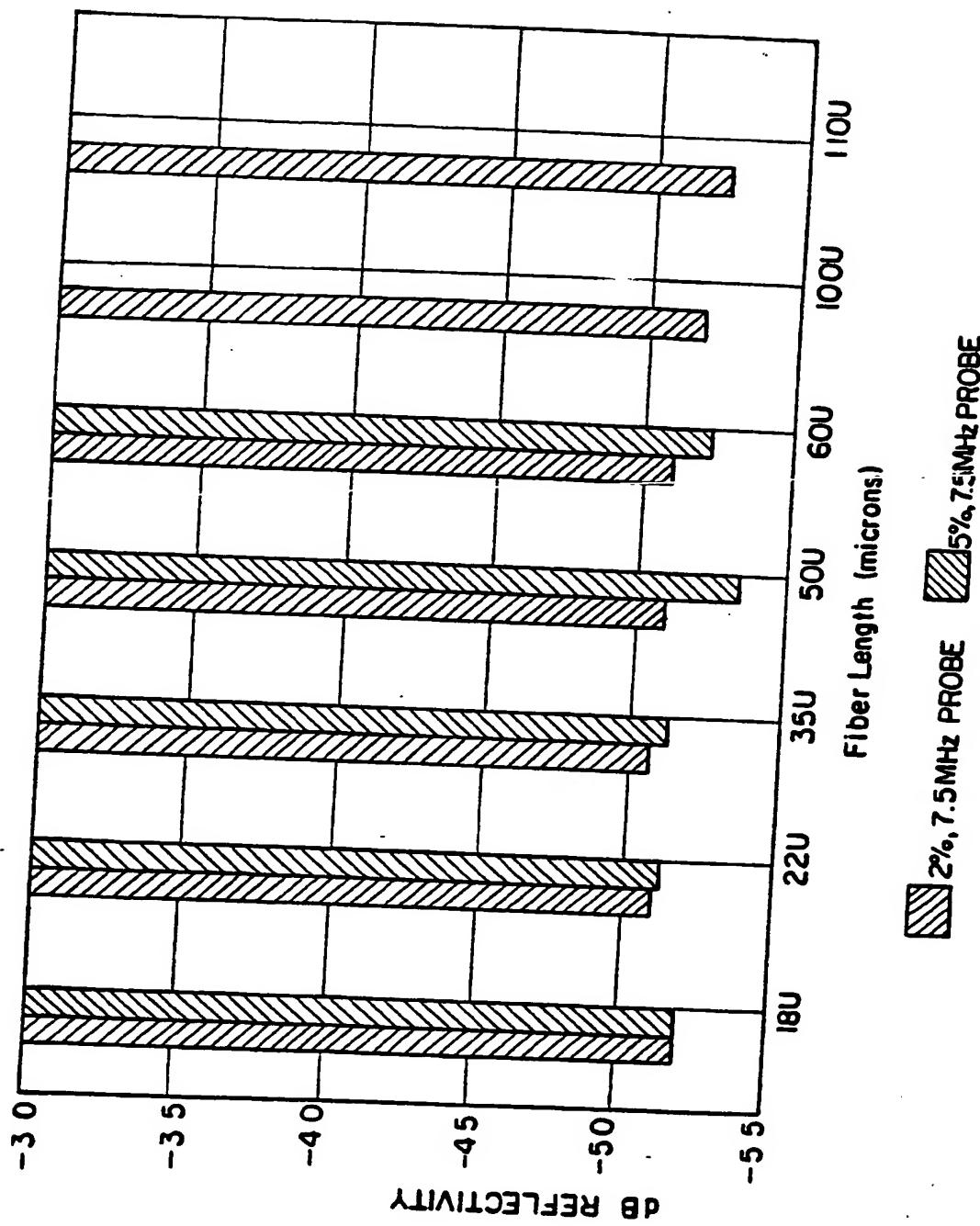


FIG. 6

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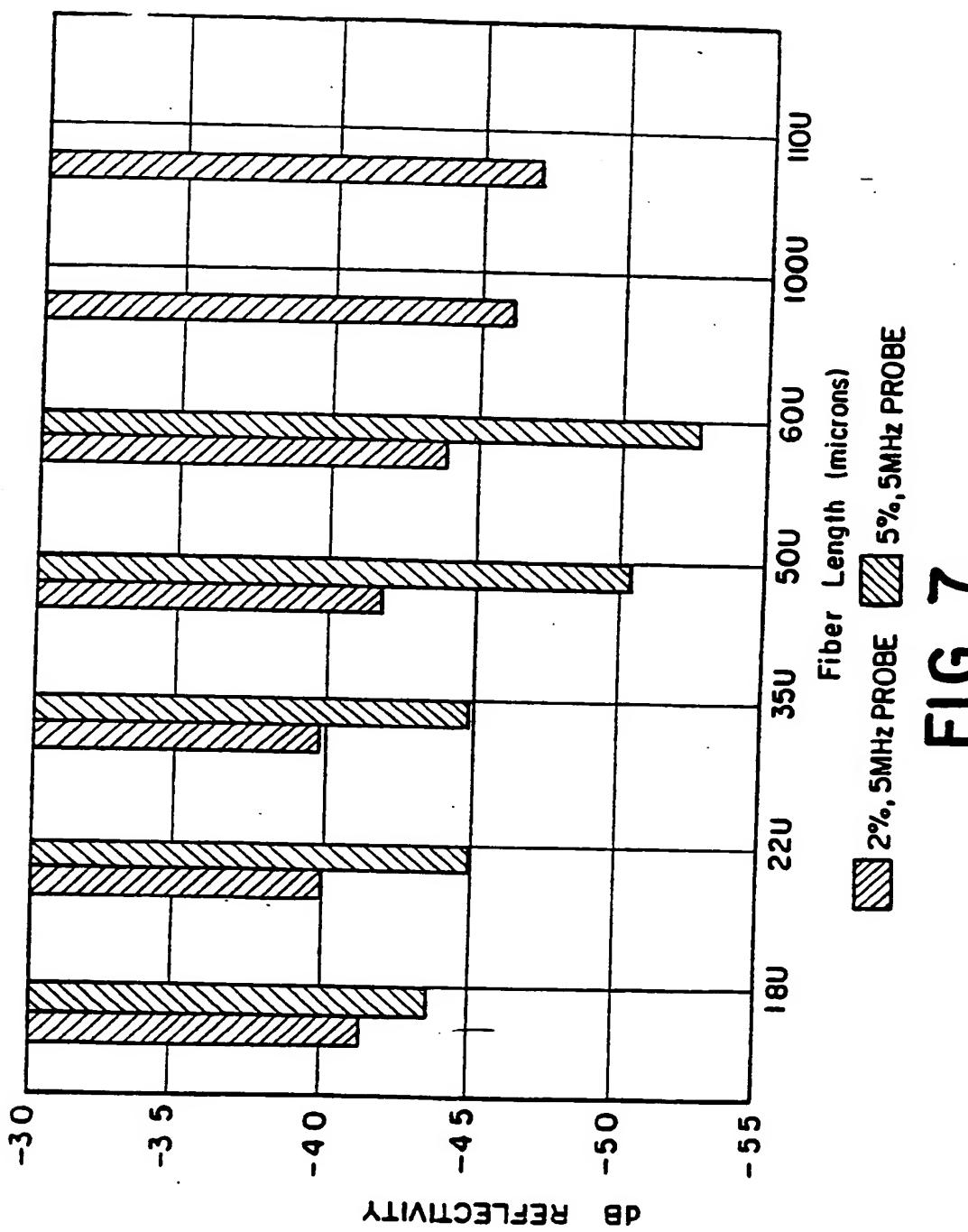
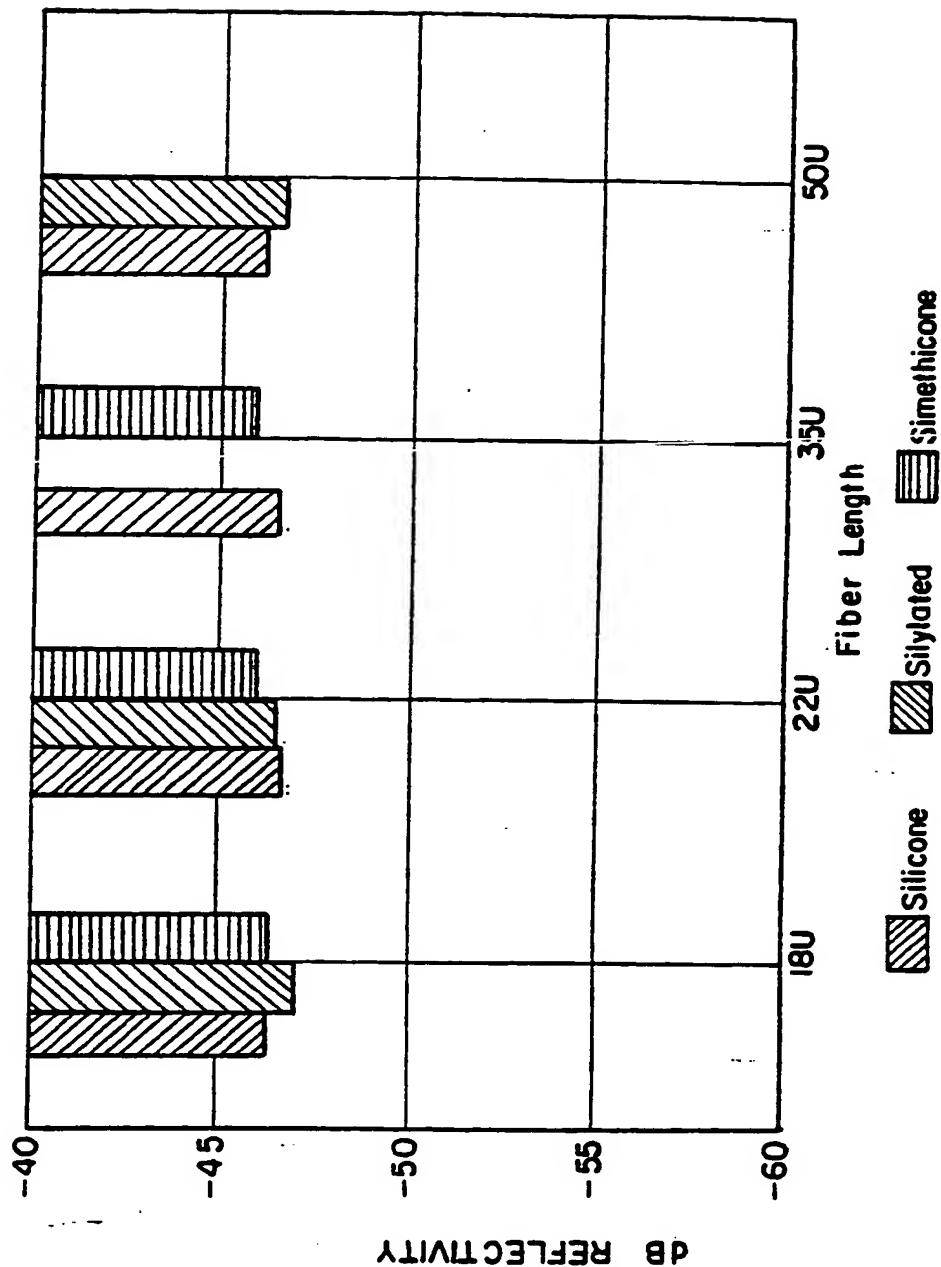


FIG. 7

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INTERNATIONAL SEARCH REPORT

International Application No.

PCT/US91/03850

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all) *

According to International Patent Classification (IPC) or to both National Classification and IPC

IPC(5): A61K 31/70, 31/715, 31/73, 31/695

U.S.C.L.: 514/54, 53, 55, 57, 58, 59, 62, 63

II. FIELDS SEARCHED

Classification System	Minimum Documentation Searched ?	
	Classification Symbols	
U.S.	514/54, 53, 55, 57, 58, 59, 62, 63	

Documentation Searched other than Minimum Documentation
to the Extent that such Documents are Included in the Fields Searched *

III. DOCUMENTS CONSIDERED TO BE RELEVANT *

Category *	Citation of Document, ¹¹ with indication, where appropriate, of the relevant passages ¹²	Relevant to Claim No. ¹³
Y	US, A, 4,101,647 (Clauss et al.) 18 July 1978, See col. 2, line 64 to col. 3, line 45.	1-40
Y	US, A, 4,269,068 (Molina) 26 May 1981, See col. 3, line 1 to col. 4, line 61.	1-29, 35-40
Y	US, A, 4,365,515 (Molina) 28 December 1982, See col. 2, lines 44 to 60.	1-29, 35-40
Y	US, A, 4,652,593 (Kawahara et al.) 24 March 1987, See col. 2, line 37 to col. 3, line 47.	1-29, 35-40
A	US, A, 4,716,225 (Ledley et al.) 29 December 1987, See entire document.	1-34

* Special categories of cited documents: ¹⁰

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

IV. CERTIFICATION

Date of the Actual Completion of the International Search

05 September 1991

Date of Mailing of this International Search Report

01 OCT 1991

International Searching Authority

ISA/US

Signature of Authorized Officer

Spruce Manis Jr

Everett White

ebw

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